



Detection of MRSA and MSSA CC398 isolates in cystic fibrosis patients of a Spanish Hospital

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Background

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Staphylococcus aureus is one of the most prevalent pathogens in cystic fibrosis (CF). (MRSA) Methicillin resistant S. aureus presence is increasingly reported, as well as the emergence of livestock-associated (LA)-MRSA lineage in CF patients.



The objective of this study was to determine the resistance phenotype/genotype, the virulence content, and molecular typing of S. *aureus* isolates from CF infections





40 S. aureus isolates were obtained from CF patients of a Spanish hospital during January-April 2022 (one isolate per patient).

multi-resistant.

PEN OXA ERY GEN TOB AMK CLO SXT CIP TET

Figure 1. Frequency of antimicrobial resistance of the 40 S. aureus isolates. PEN, penicillin; OXA, oxacillin; ERY, erythromycin; CLI, clindamycin; TET, tetracycline; GEN, gentamicin; TOB, tobramycin; AMK, amikacin; CIP, ciprofloxacin; SXT, trimethoprim-sulfamethoxazole



Figure 2. Distribution of IEC according to clonal complexes.

7 CC398 isolates were detected: 1 MRSA-t011-IEC negative 6 MSSA-t011-t034-t108-t571





The phenotype and genotype of antimicrobial resistance was evaluated by Microscan and PCR-sequencing.



The presence of lukS/lukF-PV, eta, etb, and *tst* genes was determined by PCR.



Molecular typing (agr-, spa-typing) was studied by PCR-sequencing, and the Immune Evasion Cluster (IEC) genes were analysed by PCR.

$\mathcal{A}_{\mathcal{A}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \mathcal{B}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \mathcal{B}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \mathcal{B}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \mathcal{B}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \mathcal{B}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \xrightarrow{\mathcal{B}} \xrightarrow{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \xrightarrow{\mathcal{B}}_{\mathcal{B}} \xrightarrow{\mathcal{B}$

Table 1. Characteristics of the 7 CC398 isolates of this study

Figure 3. Distribution of MRSA a MSSA according to clonal complexes.

	<i>spa</i> -type	IEC	Resistance phenotype	Resistance genotype
MRSA	t011 (1)	_	PEN, OXA, ERY, CLI, TET, GEN, TOB, AMK, CLO, CIP	blaZ, mecA, tet(K), tet(M), erm(A), aac(6')-le-aph-(2'')-la, ant(4')-la, aph(3')-III
	t011 (1)	-	PEN, ERY, CLI, TOB, AMK, CIP	<i>blaZ, erm</i> (A), <i>vga</i> (A), <i>aac</i> (6')-le- <i>aph</i> -(2'')-la
MSSA	t034 (1)	-	PEN, TET, GEN, TOB, AMK, CIP	<i>blaZ, tet</i> (M),
	t108 (2)	B ¹ /-1	PEN, ERY, CLI ¹ , TET ¹ , CIP, SXT ¹	blaZ, tet(M) ¹ , mrs(A) ¹ , mph(C) ¹
	t571 (2)	F	PEN, ERY, CLI ^{Ind} , CIP	blaZ, erm(T)

A number in superscript reflects when not all isolates of the group have the referred characteristic

PEN, penicillin; OXA, oxacillin; ERY, erythromycin; CLI, clindamycin; TET, tetracycline; GEN, gentamicin; TOB, tobramycin; AMK, amikacin; CIP, ciprofloxacin; SXT, trimethoprim-sulfamethoxazole

The MRSA isolate belonged to CC398-t011-IEC negative (animal-clade), and it was multi-resistant.

The two t571-IEC-positive-MSSA isolates presented the gene erm(T) (human-clade).

 Table 2 Percentage of occurrence of virulence genes

Clonal complaxes Virulence gene

Among the 33 non-CC398 isolates, 12 of them lacked the scn gene (36%), of the lineages CC5, CC8, CC10, CC30, CC45, CC166

The *tst* gene was detected in 13 isolates being all MSSA. The *eta* gene was detected in 2 MSSA-IEC B isolates



- In one isolate (MSSA-t130-CC330) *tst* and *eta* genes were detected at the same time.
- None isolate harboured PVL gene.



- MRSA and MSSA isolates of lineage CC398 were found in CF patients showing characteristic of both human and animal clades.
- 2. Non CC398-IEC negative were identified.
- The emergence of LA clonal lineages in CF patients should be further analysed and monitored. 3.

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